EXHIBIT 5



Pain Management pocketcard Set



General Approach to Pain Management

ASK.

Always ask patient about the presence of pain and accept the patient's report of pain.

ASSESS:

Perform a comprehensive pain assessment:

- Onset, duration, and location
- Quality (sharp, dull, diffuse, throbbing, etc)
- Intensity (1-10 scale, for example) Aggravating and alleviating factors
- Effect on function and quality of life
- Patient's goal for pain control
- Response to prior tx if condition is chronic
- History and physical examination

TREAT:

- With older adults, start dose low, go slow, but goll
- Avoid IM route, the PO route is preferred
- Treat persistent pain with regularly scheduled meds Two drugs of the same class (eq. NSAIDs) should not generally be given concurrently, however long- and short-acting opioids may be prescribed together

Avoid meperidine (per American Pain Society and ISMP) and propoxyphene (cardiotoxic and 4 efficac

MONITOR:

- Assess and reassess pain frequently
- Most opioid agonists have no analgesic ceiling dose; titrate to relief and assess for adverse effects.
- Assess, anticipate, and manage opioid adverse effects aggressively
- Discuss goals and plans with pagent and family
- Addiction rarely occurs unless there is a his of abuse
- Watch for red flags of addiction:
 - 1) Compulsive use 2) Loss of control

 3) Use despite harm

Breakthrough Pain Management General

- Use long-acting opioids around the clock for baseline management of persistent pain
- Use short-acting opioids PRN (rescue) for breakthrough pain
- · Consider using the same drug for both baseline and rescue doses whenever possible (eg long-acting morphine + short-acting morphine)

Rescue Dosing

- The rescue dose is 10%-15% of the
 - 24-h total daily dosage
 - Oral rescue doses should be available: every 1-2 h; parenteral doses every 15-30 minutes

Adjustment

- If the patient is consistently taking ≥ 3 rescue doses daily, consider increasing the baseline round-the- clock dosage
- · Recalculate rescue dose whenever the baseline dosage is changed

Example Calculate rescue dose for patient on baseline coverage of MS Contin 200 mg

- q 12.h: Calculate total daily dosage: 200 mg x 2 = 400 mg morphine/d
- 10%-15% of 400 mg = 40-60 mg short-acting morphine 3. Oral rescue dose therefore is:
 - morphine 40-60 mg PO q 1-2 h
 - Parenteral rescue dose (based on continuous infusion): Calculate based on 25%-50% of hourly dose

Pain Types Quality Type Examples Somatic nam Trauma, burns, bone metastasis Constant, sometimes throbbing or aching, tender, and localized to the site of origin Viskeral pain Renal stone passage, small bowel Poorly localized, may be referred to distant obstruction, appendicitis, cancer cutaneous site (eg, diaphragmatic irritation referred to ipsilateral shoulder), often associated with nausea or diaphoresis Neuropathic pair Nerve compression, cancer invasion Prolonged, severe, burning, lancinating, of neural structures, diabetic squeezing, hypersensitivity to pain; possible neuropathy, postherpetic neuropathy, trigeminal neuralgia tachycardia, diaphoresis; tends to be resistant to opioids and difficult to treat Interventional Pain Management Techniques Technique

| Lumbar epidural steroid injection (LESI) | Inflammation associated with conditions such as spinal stenosis, disc herniation or degenerative disc disease |
|---|--|
| Facel block | Diagnostic tool used to isolate and confirm the specific source of back pain (facet joints) |
| Selective nerve root block (SNRB) | Primarily used to diagnose the specific source of nerve root pain and, secondarily, for therapeutic purposes such as treatment for a far lateral |

disc herniation Neurolytic blocks (chemical, Good for localized pain not requiring multiple segmental blocks; radiofrequency abiation) successful SNRBs should be done prior to neurolysis

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Pain Treatment Ladder

Severe-Intractoble Pai

Strong (V/Immeth ceal/intespinal counts (+/- aminor ts)
 Three procedures (merce block, meetid only, nerve/spinal stimulation).

Moderate-Severe Pain:

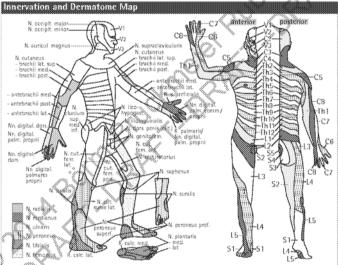
1. Weak PO opiods/opiod combination drugs

Strong PO/IV opiods if 1 ineffective; consider using adjuvants, especially for neuropathic pain

Mild-Moderate Pain:

 Nonopioid analgesics = 2. Weak PO opioids/opioid combination drugs, if 1 ineffective Mild Pain: Nonopioid analgesics

| erm | Definition | Term | Definition |
|----------------------|--|---------------|--|
| Allodynia | Feeling nonpainful stimulation as painful | Hyperpathia | Reduced sensation |
| nalgesia | No pain | Hypoesthesia | Decreased cutaneous stimulation |
| nesthesia | No sensation | Paresthesia | Abnormal sensation without stimulu |
| nesthesia olorosa | Pain in an area with no stimulation | Hyperesthesia | Increased response to muldistimuli |
| lypoalgesia | Diminished response to pain | Dysesthesia | Unpleasant sensation with or withou stimulation. |



| Drug | Onset | Duration (h) | CNS Tox | Heart Tox | Pot | Comments |
|------------------------|-------|-------------------------------------|---------|-----------|---------------------------|--|
| Amides | | | | | | Slow, hepatic metabolism; |
| idocaine | fast | 1-2 | ++ | + | 4 | high systemic toxicity potential, but low allergic |
| Supivacaine | slow | 3-6 | +++ | ++++++ | 16 | potential; bupivacaine has |
| depivacaine | mod | 1-3 | ++ | + | 3-4 | high cardiotoxic potential; prilocaine is associated with methemoglobinemia at hig doses |
| ^a rilocaine | fast | 2-3 | + | +/- | 3-4 | |
| lopvacaine | mod | Epidural ~7 PNB ² 2-6 | ++{+} | +++ | 16 | |
| Esters | | | | | Rapid metabolism by plasn | |
| rocaine | fast | 0.5-1 | + | + | 1 | cholinesterase; high allergi- potential (PABA derivatives tetracaine is the most toxic among the esters |
| htoroprocaine | fast | 0.5-1 | + | + | 4 | |
| etracaine | slow | 1.5-3 | +++ | +++ | 16 | |

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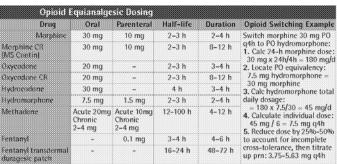












| Opioid Patient-C | Controlled Ana | lgesia (PCA) Ri | egimens ^l | |
|------------------|----------------|-----------------|------------------------|------------------|
| Drug | Conc. [mg/mt] | Bolus dose (mg) | Lockout interval (min) | Hourly max [stg] |
| Morphine | 1 | 0.5-2.5 | 5-15 | §-15 |
| Fentanyl | 0.025 or 0.050 | 0.0125-0.050 | 5-10 | 0.1-0.3 |
| Hydromorphone | 1 | 0.3 | 5-15 | 1.25-3 |

These dosages are for opioid-naive patients; much larger dosages may be needed for opioid-tollerant patients.

| Drug | Dose | Adverse Effects | Comments |
|---------------------------------|--|---|---|
| Codeme | 15-120 mg PO/IM/ SC q 4-6 h | Drowsiness, constipation, bradycardia, euphoria, confusion, pruritus | Requires dosage reduction in renal failure |
| Iramadel | 25-50 mg q 4-6 h Max 400 mg/d, 300 mg/d in elderly | Headache, conflision, sedation | Dual-action opioid agonist, norepi/serotonin receptor antagonist; ↓ seizure threshold |
| Hydrocorlone + scetaminophen | 1 tab (2.5–10 mg / 325–750 mg) PO q 4–6 h prn | Sedation, respiratory depression, hypotension, pruritus, confusion, constipation | Max 4 g/day acetaminophen |
| Oxycodone i acetaminophen | 1 tab (2.5–10mg / 300–650 mg) PO q 4–6 h prn | Similar opioid effects | Max 4 g/day acetaminophen |

| Nonopioid Anal | gesies | | |
|---|---|--|--|
| Drug | Dostige | Adverse Effects | Comments |
| NSAIDs /* | | | |
| Aspirin | 800-1000 mg g4-6h Max 4 g/d | GI bleeding, ↓ platelet adhesiveness, renal toxicity | Caution in hepatic/renal disease |
| Choline magiecitum trisalicylate (Infisale) | 500 mg initial then 250 mg q 6-8 h Max 1500 mg/d | Lower incidence of GI effects | Caution in hepatic/renal disease; does not inhibit platelet aggregation |
| lhuprofén | 200–400 mg q 4–6 h Max 2400 mg/d | GI bleeding, ↓ platelet adhesiveness, renal toxicity | Caution in hepatic/renal disease |
| Maproxen | 500 mg initial then 250 mg q 6-8 h Max 1500 mg/d | GI bleeding, ‡ platelet adhesiveness, renal toxicity | Caution in hepatic/renal disease |
| Naburbetone | 500-750 mg q8-12h Max 2 g/d | GI bleeding, ↓ platelet adhesiveness, renal toxicity | Caution in hepatic/renal disease |
| Ketomise | 30 mg Vinitial, then 15–30 mg q 6 h Max 150 mg/d day 1, then 120 mg/d | GI bleeding, i platelet adhesiveness, renal toxicity | In elderly 30 mg IV initial, then 15-30 mg thereafter. Use restricted to max 5 days. Caution in hepatic/renal disease |
| Celecoxib | 100-200 mg q 12 h Max 200-400 mg/d | Lower incidence of GI effects | Does not inhibit platelet aggregation |
| Other | | | |
| Acetaminophen | 500-1000 mg q4-6h Max 4 g/d, 3 g/d if liver dis or elderly | Liver toxicity at high doses | Use caution in the elderly and individuals with hepatic disease |
| Ziconotide | µg/d; titrate by ≤2.4 | Neurologic and cognitive impairment, dizziness, confusion, memory deficits, NIV/D. 1 CK | N-type Ca channel blocker; for intractable pain unresponsive to other agents |

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| Analoesic Adjuv Drug | Dosage | Adverse Effects | Comments |
|---------------------------------|---|---|--|
| Antidepressants ² | | | |
| Amitriplyline | Init 25 mg PO qhs Increase to 100 mg PO qhs prn | Sedation, constipation, urinary retention, tachycardia, conduction abnormalities, seizures | Tricyclic antidepressant (TCA); has the most anticholinergic effects |
| Desipramine | 100 mg PO qd | similar effects | TCA; fewer adverse effects |
| Imipramine | 100 mg PO qd | similar effects | TCA |
| Nortriptyline | 50-100 mg PO qhs | similar effects | TCA; less sedating |
| Duloxetine | 60 mg PO qd Max 120 mg/d | Sedation, insomnia, dizziness, nausea | SNRI; indicated for diabetic neuropathic pain |
| Anticonvulsants | | | |
| Carbamazepine | Init 100 mg PO bid Titrate to max of 1,600 mg/d div qid | Nausea, vomiting, diarrhea, hyponatremia, rash, pruritus, drowsiness, blurred vision, headache, dizziness, Stevens-Johnson syndrome | Indicated for trigeminal or glossopharyngeal neuralgia; requires CBC and LFT monitoring; Asians with the HLA-B*1502 allele are predisposed to Stevens 388psor |
| Gabapentin | Day 1: 300mg PO qhs Day 2: 300 mg PO bid Day 3: 300 mg PO tid Max 1,800 mg/d PO div tid | | Indicated for postherpetic neuralgia; requires dose reduction in cenal, failure |
| Pregabalin | Init 50 mg PO tid Max 100 mg PO tid | Weight gain, somnolence, dizziness, ataxia, peripheral edema | Indicated for postherpetic negration, diabetics neuropathic pain, fibromyalgia; requires dose reduction in renal failure |
| Other Agents Capsaicin cream | 0.025%-0.075% | Itching, stinging, erythema | |
| Lidocaine 5% patch | Up to 3 patches at once for up to 12 h within 24 h period | Local skin reactions such as blisters or erythema | Indicated for postherpetic neuralgia |
| Clonidine | Epidural infusion as opiate adjunct: init 0.5 μg/kg/h; † dose to effect | Drowsiness, chrziness, dry mouth, constitution, skin reactions, orthostatic hypotension | Opiate adjunct for severe, intractable pain, unresp to other analgesics or spinal opiates alone, esp neuropathic pain |

May be used alone or in combination with epioids, often in the treatment of neuropathic pain.

With the exception of duloxetine, use of these agents in pain management is off-label, however, they are considered by pain specialists as first-line treatment in diabetic peripheral neuropathy pain (DPNP).

Anticoagulation in Neuraxial Ancethesia Orug Minimum Elapsed Time Drug Minimum Elapsed Time Delay for 1 h after needle placement; ASA/NSAIDs No risk Witeparin remove indwelling catheters 2-4 h after last dose Clopidogret 7 days Abciximab 48 h 4-5 days Entifibatide 8 h Warfarin 14 days E NAMAZEA 10-12 h (low dose); 24 h (high dose) Fictopidine SubO Heparin No risk Innombolytics Avoiding regional block is recommended Minimum clapsed time between the last drug dose and administration of anesthesia

Management of Opioid Adverse Events

Adverse Event Management Constitution Begin bowel regimen when opioid therapy is initiated. Include a mild stimulant laxative (eg, Senna, Cascara) + stool softener (eg, Colace) at bedtime or in divided doses as routine prophylaxis.

Sedation Tolerance typically develops. Hold sedatives/anxiolytics, reduce opioid dose. Consider stimulants such as caffeine, methylphenidate, or dextroamphetamine. Naysca/vomiting Dosage reduction, opioid rotation. Consider transdermal scopolamine patch, metoclopramide, or prochlorperazine. Caused by opioid induction of histamine release that is inversely correlated to

potency (morphine > fentanyl). Management involves dosage reduction, opioid rotation, and possible use of an antihistamine (eg, diphenhydramine) Hallucinations Dosage reduction, opioid rotation. Consider neuroleptics [eg, haloperidol, risperidone] Confusion/delirium Dosage reduction, opioid rotation, neuroleptic therapy (eg, haloperidol, risperidone) Myoclonic jerking Dosage reduction, opioid rotation. Consider clonazepam, baclofen. Sedation precedes respiratory depression. Stop opioid! Give low-dose naloxone Respiratory dilute 0.4 mg (1 mL of a 0.4 mg/mL amp of naloxone) in 9 mL of normal saline (NS)

for final concentration of 0.04 mg/mL. Recommendations for Treatment of Diabetic Peripheral Neuropathy Pain (DPNP) ist-tier drugs Duloxetine, oxycodone CR, pregabalin, tricyclic antidepressant (ICA) class drugs 2nd-tier drugs Carbamazepine, gabapentin, lamotrigine, tramadol, venlafaxine Honorable mention: Topical capsaicin, topical lidocaine, bupropion, citalopram, paroxetine, phenytoin, topiramate, methadone

Adapted from the Mayo Clinic 2006 Consensus Guidelines for the treatment of DPNP.

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Pruritus

depression





